## **AMENDMENTS TO THE CLAIMS**

Docket No.: 12810-00276-US1

## **Listing of Claims**

- 1. (Currently Amended) A method of separating acids from reaction mixtures by means comprising the use of an auxiliary base, where the auxiliary base
  - A) reacts with the acid to form a salt which is liquid at temperatures at which the desired product is not significantly decomposed while the liquid salt is being separated off and
  - B) the salt of the auxiliary base forms two immiscible liquid phases with the desired product or the solution of the desired product in a suitable solvent,

wherein and wherein the method the auxiliary base used is an alkylimidazole

- which has a solubility in 30% strength by weight sodium chloride solution at 25°C of 10% by weight or less and
- whose hydrochloride has a melting point below 55°C;

which comprises the following steps:

- a) reacting at least one 1-alkylimidazole with at least one acid in the presence of a desired product to form a mixture of at least one salt of the 1-alkylimidazole and the desired product,
- b) separating the salt or salts of the 1-alkylimidazole and the desired product under conditions under which at least two separate phases of which at least one comprises predominantly the salt or salts of the 1-alkylimidazole and at least one other comprises predominantly desired product are formed,
- c) adding at least one base to a phase which has been separated off from (b) and comprises predominantly the salt or salts of the 1-alkylimidazole to form a mixture of the liberated 1-alkylimidazole and the reaction product of base and acid,
- d) separating the mixture of the liberated 1-alkylimidazole and the reaction product of base and acid under conditions under which at least two separate phases of which at least one comprises predominantly the liberated 1-alkylimidazole in crude form and at least one other comprises the reaction product of base and acid are formed,
- e) if appropriate, purifying the 1-alkylimidazole obtained in crude form and

f) if appropriate, recirculating the optionally purified 1-alkylimidazole to step (a).

Docket No.: 12810-00276-US1

- 2. (Original) The method according to claim 1, wherein a 1-alkylimidazole whose hydrochloride has a melting point below 45°C is used.
- 3. (Currently Amended) The method according to claim 1 or 2 claim 1, wherein a 1-alkylimidazole having a solubility in 30% strength by weight sodium chloride solution at 25°C of 3% by weight or less is used.
- 4. (Currently Amended) The method according to any of claims 1 to 3 claim 1, wherein the auxiliary base used is a 1-alkylimidazole of the formula (I),

$$\begin{array}{c|c}
 & H_2 \\
 & C \\
 & C \\
 & C \\
 & R^1
\end{array}$$
(I)

where  $R^1$  and  $R^2$  can each be, independently of one another, hydrogen or linear or branched  $C_1 - C_6$ -alkyl, with the proviso that  $R^1$  and  $R^2$  have a total of at least 1 carbon atom and a total of not more than 6 carbon atoms.

- 5. (Original) The method according to claim 4, wherein R<sup>1</sup> and R<sup>2</sup> are selected independently from the group consisting of hydrogen, methyl and ethyl.
- 6. (Currently Amended) The method according to any of the preceding claims claim 1, wherein the 1-alkylimidazole is selected from the group consisting of 1-n-propylimidazole, 1-n-butylimidazole and 1-isobutylimidazole.
- 7. (Original) The method according to claim 6, wherein the separation of the phases in step (b) is carried out in a phase separator.
- 8. (Currently Amended) The method according to claim 1 or 7 claim 1, wherein the concentration of the base or bases added in step (c) is selected so that the reaction product of base and acid in step (d) is obtained in at least 15% strength by weight solution.

9. (Currently Amended) The method according to any of claims 1, 7 or 8 claim 1, wherein the purification in step (e) comprises single or multiple washing, drying, filtration, stripping, distillation and/or rectification.

Docket No.: 12810-00276-US1

- 10. (New) The method according to claim 2, wherein a 1-alkylimidazole having a solubility in 30% strength by weight sodium chloride solution at 25°C of 3% by weight or less is used.
- 11. (New) The method according to claim 2, wherein the auxiliary base used is a 1-alkylimidazole of the formula (I),

$$\begin{array}{c|c} & H_2 \\ & C \\ & C \\ & C \\ & R^1 \end{array}$$

where  $R^1$  and  $R^2$  can each be, independently of one another, hydrogen or linear or branched  $C_1$  –  $C_6$ -alkyl, with the proviso that  $R^1$  and  $R^2$  have a total of at least 1 carbon atom and a total of not more than 6 carbon atoms.

12. (New) The method according to claim 3, wherein the auxiliary base used is a 1-alkylimidazole of the formula (I),

$$\begin{array}{c|c}
 & H_2 \\
 & C \\
 & C \\
 & C \\
 & R^1
\end{array}$$
(I)

where  $R^1$  and  $R^2$  can each be, independently of one another, hydrogen or linear or branched  $C_1$  –  $C_6$ -alkyl, with the proviso that  $R^1$  and  $R^2$  have a total of at least 1 carbon atom and a total of not more than 6 carbon atoms.

13. (New) The method according to claim 2, wherein the 1-alkylimidazole is selected from the group consisting of 1-n-propylimidazole, 1-n-butylimidazole and 1-isobutylimidazole.

Application No. National Phase of PCT/EP04/014386 Amendment dated June 16, 2006 First Preliminary Amendment

14. (New) The method according to claim 3, wherein the 1-alkylimidazole is selected from the group consisting of 1-n-propylimidazole, 1-n-butylimidazole and 1-isobutylimidazole.

Docket No.: 12810-00276-US1

15. (New) The method according to claim 4, wherein the 1-alkylimidazole is selected from

the group consisting of 1-n-propylimidazole, 1-n-butylimidazole and 1-isobutylimidazole.

16. (New) The method according to claim 5, wherein the 1-alkylimidazole is selected from

the group consisting of 1-n-propylimidazole, 1-n-butylimidazole and 1-isobutylimidazole.

17. (New) The method according to claim 2, wherein the concentration of the base or bases

added in step (c) is selected so that the reaction product of base and acid in step (d) is obtained in

at least 15% strength by weight solution.

18. (New) The method according to claim 3, wherein the concentration of the base or bases

added in step (c) is selected so that the reaction product of base and acid in step (d) is obtained in

at least 15% strength by weight solution.

19. (New) The method according to claim 4, wherein the concentration of the base or bases

added in step (c) is selected so that the reaction product of base and acid in step (d) is obtained in

at least 15% strength by weight solution.

20. (New) The method according to claim 5, wherein the concentration of the base or bases

added in step (c) is selected so that the reaction product of base and acid in step (d) is obtained in

at least 15% strength by weight solution.

6